PALM INTRANET

Day : Thursday
Date: 9/5/2002
Time: 15:38:14

Inventor Name Search Result

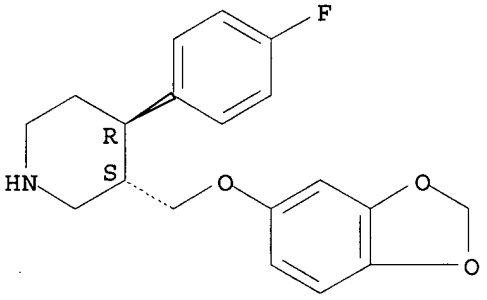
Your Search was:

Last Name = BENNEKER
First Name = FRANCISCUS

Application#	Patent#	Status	Date Filed	Title	Inventor Name
09428529	Not Issued	061	10/28/1999	PROCESS FOR PRODUCING 4-ARYLIPIPERIDINE-3-CARBINOLS AND RELATED COMPOUNDS	BENNEKER , FRANCISCUS B.G.
60106673	Not Issued	159	11/02/1998	PROCESS FOR PRODUCING 4-ARYPIPERIDINE-3-CARBINOLS AND RELATED COMPOUNDS	BENNEKER , FRANCISCUS B.G.
08872023	5874447	150	06/10/1997	4-PHENYLPIPERIDINE COMPOUNDS FOR TREATING DEPRESSION	BENNEKER , FRANCISCUS BERNARDUS G
09200743	Not Issued	172	11/30/1998	4-PHENYLPIPERIDINE COMPOUNDS	BENNEKER , FRANCISCUS BERNARDUS GEMM
09938840	Not Issued	071	08/27/2001	PROCESS FOR MAKING AMLODIPINE, DERIVATIVES THEROF, AND PRECURSORS THEREFOR	BENNEKER, FRANCISCUS B. G.
60258602	Not Issued	020	12/29/2000	ASPARTATE DERIVATIVE OF AMLODIPINE	BENNEKER, FRANCISCUS B. G.
09938818	Not Issued	041	08/27/2001	AMIDE DERIVATIVE OF AMLODIPINE	BENNEKER, FRANCISCUS B. G.
09938817	Not Issued	093	08/27/2001	ASPARTATE DERIVATIVE OF AMLODIPINE	BENNEKER, FRANCISCUS B.G.
09938841	Not Issued	071	08/27/2001	PROCESS FOR MAKING AMLODIPINE MALEATE	BENNEKER, FRANCISCUS B.G.
10024520	Not Issued	041	12/21/2001	AMLODIPINE FREE BASE	BENNEKER, FRANCISCUS B.G.

RN 78246-49-8 REGISTRY
CN Piperidine, 3-[(1,3-benzodioxol-5-yloxy)methyl]-4-(4-fluorophenyl)-, hydrochloride, (3S,4R)- (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN Piperidine, 3-[(1,3-benzodioxol-5-yloxy)methyl]-4-(4-fluorophenyl)-, hydrochloride, (3S-trans)-
OTHER NAMES:
CN (-)-trans-4-(4-Fluorophenyl)-3-(3,4-methylenedioxyphenoxymethyl)piperidine hydrochloride
CN Aropax 20
CN BRL 29060 hydrochloride
CN BRL 29060A
CN Deroxat
CN Paroxet
CN **Paroxetine hydrochloride**
CN Paxil
CN Seroxat
CN Tagonis
FS STEREOSEARCH
DR 172501-13-2
MF C19 H20 F N O3 . Cl H
CI COM
LC STN Files: BIOBUSINESS, BIOSIS, CA, CAPLUS, CASREACT, CHEMCATS, CIN, CSCHEM, DIOGENES, DRUGPAT, DRUGUPDATES, IPA, MRCK*, PHARMASEARCH, PROMT, RTECS*, TOXCENTER, USPATFULL
(*File contains numerically searchable property data)
CRN (61869-08-7)

Absolute stereochemistry. Rotation (-).



O HCl

108 REFERENCES IN FILE CA (1967 TO DATE)
2 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
109 REFERENCES IN FILE CAPLUS (1967 TO DATE)

L2 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2002 ACS
RN 78246-49-8 REGISTRY
CN Piperidine, 3-[(1,3-benzodioxol-5-yloxy)methyl]-4-(4-fluorophenyl)-, hydrochloride, (3S,4R)- (9CI) (CA INDEX NAME)
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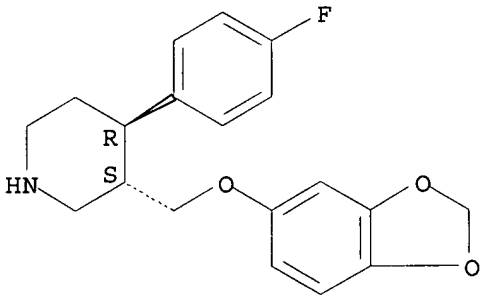
OTHER NAMES:
CN (-)-trans-4-(4-Fluorophenyl)-3-(3,4-methylenedioxyphenoxymethyl)piperidine hydrochloride
CN Aropax 20
CN BRL 29060 hydrochloride
CN BRL 29060A
CN Deroxat
CN Paroxet
CN **Paroxetine hydrochloride**
CN Paxil
CN Seroxat
CN Tagonis
FS STEREOSEARCH
DR 172501-13-2
MF C19 H20 F N O3 . Cl H
CI COM
LC STN Files: BIOBUSINESS, BIOSIS, CA, CAPLUS, CASREACT, CHEMCATS, CIN, CSCHEM, DIOGENES, DRUGPAT, DRUGUPDATES, IPA, MRCK*, PHARMASEARCH, PROMT, RTECS*, TOXCENTER, USPATFULL
(*File contains numerically searchable property data)
CRN (61869-08-7)

Absolute stereochemistry. Rotation (-).



O HCl

108 REFERENCES IN FILE CA (1967 TO DATE)
2 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
109 REFERENCES IN FILE CAPLUS (1967 TO DATE)

=> fil caplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	5.96	34.47

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION

CA SUBSCRIBER PRICE 0.00 -5.58

FILE 'CAPLUS' ENTERED AT 10:42:18 ON 29 AUG 2002
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FILE COVERS 1907 - 29 Aug 2002 VOL 137 ISS 9
FILE LAST UPDATED: 27 Aug 2002 (20020827/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

CAS roles have been modified effective December 16, 2001. Please check your SDI profiles to see if they need to be revised. For information on CAS roles, enter HELP ROLES at an arrow prompt or use the CAS Roles thesaurus (/RL field) in this file.

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=> s l2/p
L3          46 L2/P

=> s l3(l)(acetate or maleate)
        407426 ACETATE
         26135 MALEATE
L4           3 L3(L)(ACETATE OR MALEATE)

=> d bib abs 1-3
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L4 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2002 ACS
AN 2000:457064 CAPLUS
DN 133:73944
TI Salification process for the preparation of an acetate salt of paroxetine or paroxetine analogs
IN Craig, Andrew Simon; Jones, David Alan; Man, John
PA Smithkline Beecham PLC, UK
SO PCT Int. Appl., 15 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000039123	A1	20000706	WO 1999-GB4370	19991222

W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

EP 1140912 A1 20011010 EP 1999-962415 19991222
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO
PRAI GB 1998-28781 A 19981229
WO 1999-GB4370 W 19991222
OS CASREACT 133:73944; MARPAT 133:73944
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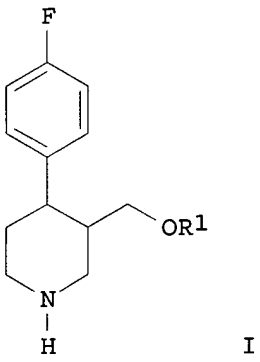


I

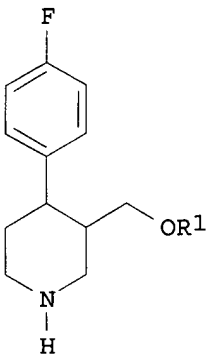
AB Acetate salts of paroxetine and its analogs (I; R1 = substituted Ph, preferably 3,4-methylenedioxyphenyl) (e.g., paroxetine acetate), useful as therapeutic agents (no data), are prepd. by contacting a soln. of the I (e.g., paroxetine) base with an amine-acetic acid salt (e.g., ammonium acetate).

L4 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2002 ACS
AN 2000:457063 CAPLUS
DN 133:73943
TI Process for the preparation of an acetate salt of paroxetine or paroxetine analogues
IN Craig, Andrew Simon; Jones, David Alan; Man, John
PA Smithkline Beecham Plc, UK
SO PCT Int. Appl., 15 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2000039122	A1	20000706	WO 1999-GB4367	19991222
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
EP 1140911	A1	20011010	EP 1999-962412	19991222
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				

PRAI GB 1998-28780 A 19981229
WO 1999-GB4367 W 19991222
OS CASREACT 133:73943; MARPAT 133:73943

GI



I

AB A process for the prepn. of an acetate salt of paroxetine or an analog. (I; R1 = substituted Ph group, preferably 3,4-methylenedioxyphenyl) (e.g., paroxetine acetate) comprises crystg. the acetate salt from a solvent (e.g., toluene) capable of dehydrating an aq. soln. by forming an azeotrope.

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2002 ACS
AN 2000:457062 CAPLUS
DN 133:79375
TI Preparation of an acetate salt of paroxetine or its analogs
IN Craig, Andrew Simon; Jones, David Alan; Man, John
PA Smithkline Beecham P.L.C., UK
SO PCT Int. Appl., 16 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000039121	A1	20000706	WO 1999-GB4364	19991222
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	EP 1140910	A1	20011010	EP 1999-962409	19991222
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
PRAI	GB 1998-28779	A	19981229		
	WO 1999-GB4364	W	19991222		

OS MARPAT 133:79375
AB The present invention relates to a new process for the prepn. of an acetate salt of a compd. paroxetine or its analogs. The (-) isomer of paroxetine has antidepressant and anti-Parkinson properties. This compd. is used in therapy as a hydrochloride salt to treat inter-alia depression obsessive compulsive disorder (OCD) and panic. A soln. of paroxetine base

in toluene, which had previously been dried over anhyd. magnesium sulfate, was dild. with propan-2-ol and seeded with cryst. paroxetine acetate. Acetic acid was added and the soln. stirred at 20.degree. overnight. The resulting solid was filtered to give paroxetine acetate as a white cryst. solid.

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT